

Prisca 5.1.0.17
Date of report: 19-07-2025

N A

Patient data			
Name	Mrs. A MAHIMA	Patient ID	0012507160268
Birthday	10-09-2005	Sample ID	B3108119
Age at sample date	19.8	Sample Date	15-07-2025
Gestational age	13 + 3		
Correction factors			
Fetuses	1	IVF	no
Weight	46	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	unknown
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age
PAPP-A	2.29 mIU/mL	0.30	13 + 2
fb-hCG	33.74 ng/mL	0.93	Method
			CRL Robinson
			Scan date
			14-07-2025
Risks at sampling date			Trisomy 21
Age risk		1:1123	The calculated risk for Trisomy 21 (with nuchal translucency) is below the cut off, which indicates a low risk.
Biochemical T21 risk		1:369	After the result of the Trisomy 21 test (with NT) it is expected that among 2497 women with the same data, there is one woman with a trisomy 21 pregnancy and 2496 women with not affected pregnancies.
Combined trisomy 21 risk		1:2497	The PAPP-A level is low.
Trisomy 13/18 + NT		<1:10000	The calculated risk by PRISCA depends on the accuracy of the information provided by the referring physician.
			Please note that risk calculations are statistical approaches and have no diagnostic value!
			The patient combined risk presumes the NT measurement was done according to accepted guidelines (Prenat Diagn 18: 511-523 (1998)).
			The laboratory can not be hold responsible for their impact on the risk assessment ! Calculated risks have no diagnostic value!
Trisomy 13/18 + NT			
The calculated risk for trisomy 13/18 (with nuchal translucency) is < 1:10000, which represents a low risk.			

Sign of Physician